

12-18-06

Attorney Docket No. 57070-8021.US00

SFW  
1625



PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLICATION OF: YUAN et al.  
APPLICATION No.: 10/591,358  
FILED: August 31, 2006  
FOR: TRIPTOLIDE DERIVATIVES FOR  
MDULATION OF APOPTOSIS AND  
IMMUNOSUPPRESSION

EXAMINER: To be Assigned  
ART UNIT: 1625  
CONF. NO: 9546

Information Disclosure Statement Within Three Months of Application  
Filing or Before First Action – 37 C.F.R. § 1.97(b)

Mail Stop Amendment  
Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

Sir:

1. Timing of Submission

This information disclosure is being filed within three months of the filing date of this application or date of entry into the national stage of an international application or before the mailing date of a first Office action on the merits, whichever occurs last (37 C.F.R. § 1.97(b)). The references listed on the enclosed Form PTO-1449 (modified) may be material to the examination of this application; the Examiner is requested to make them of record in the application.

2. Cited Information

- Copies of references 1-8 are issued patent(s) and published application(s) and are not included (see C.F.R. § 1.98(a)(2)(i)).
- Copies of references 9-28 are enclosed.

3. Effect of Information Disclosure Statement (37 C.F.R. § 1.97(h))

This Information Disclosure Statement is not to be construed as a representation that: (i) a search has been made; (ii) additional information material to the examination of this application does not exist; (iii) the information, protocols, results and the like reported by third parties are accurate or enabling; or (iv) the cited information is, or is considered to be, material to patentability. In addition, applicant does not admit that any enclosed item of information constitutes prior

art to the subject invention and specifically reserves the right to demonstrate that any such reference is not prior art.

4. Fee Payment

No fees are believed due because this Information Disclosure Statement is being filed before the mailing date of the first Office Action.

However, should the Commissioner determine that fees are due in order for this Information Disclosure Statement to be considered, the Commissioner is hereby authorized to charge such fees to Deposit Account No. 50-2207.

5. Patent Term Adjustment (37 C.F.R. § 1.704(d))

The undersigned states that each item of information submitted herewith was cited in a communication from a foreign patent office in a counterpart application and that this communication was not received by any individual designated in 37 C.F.R. § 1.56(c) more than thirty days prior to the filing of this statement. 37 C.F.R. § 1.704(d).

Respectfully submitted,  
Perkins Coie LLP

Date: December 14, 202X



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**INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT**  
Form PTO-1449 (Modified)  
(Use several sheets if necessary)

Sheet 1 of 2 Attorney Docket No. 57070-8021.US00

**COMPLETE IF KNOWN**

Application Number	10/591,358
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First Named Inventor	Yuan et al.
Group Art Unit	1625
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Examiner Name	To be Assigned

**U.S. PATENT DOCUMENTS**

Examiner Initials	Cite No.	U.S. Patent or Application		Name of Patentee or Inventor of Cited Document	Date of Publication or Filing Date of Cited Document	Pages, Columns, Lines, Where Relevant Figures Appear
		NUMBER	Kind Code (if known)			
/N.C./	4,005,108			Kupchan et al.	7/87	
	2. 5,294,443			Lipsky et al.	3/94	
	3. 5,663,335	A		Qi et al.	9/97	
	4. 5,962,516	A		Qi et al.	10/99	
	5. 5,972,998			Jung et al.	10/99	
	6. 6,004,999	A		Jung et al.	12/99	
	7. 6,150,539	A		Musser	11/00	
/N.C./	6,569,893	B2		Dai et al.	5/03	

**FOREIGN PATENT DOCUMENTS**

Examiner Initial	Cite No.	Foreign Patent or Application			Name of Patentee or Applicant of Cited Document	Date of Publication or Filing Date of Cited Document	Pages, Columns, Lines, Where Relevant Figures Appear	T
		Office	NUMBER	Kind Code				
/N.C./	9. JP	03 178977			Chugoku Igaku Kagak	8/91		
	10. PCT	WO00/12483			Pharmagenesis, Inc.	3/00		
	11. PCT	WO97/31920			Pharmagenesis, Inc.	9/97		
	12. PCT	WO97/31921			Pharmagenesis, Inc.	9/97		
	13. PCT	WO98/52933			Hoechst Marion Roussel Inc.	11/98		
	14. /N.C./	WO98/52951			Hoechst Marion Roussel Inc.	11/98		

**OTHER PRIOR ART-NON PATENT LITERATURE DOCUMENTS**

Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume/issue number(s), publisher, city and/or country where published.	T
/N.C./	15.	Anderson et al., "Synthesis, Evaluation of Chemical Reactivity, and Murine Antineoplastic Activity of 2-Hydroxy-5-(3,4-dichlorophenyl)-6,7-bis(hydroxymethyl)-2,3-dihydro-1H-pyrolizine Bis(2-propylcarbamate) and 2-Acyloxy Derivatives as Potential Water-Soluble Prodrugs <sup>1", J. Med. Chem., 26:1333-1338 (1983).</sup>	
/N.C./	16.	de Groot Franciscus M. H. et al., "Synthesis and Biological Evaluation of 2'-Carbamate-Linked and 2'-Carbonate-Linked Prodrugs of Paclitaxel: Selective Activation by the Tumor-Associated Protease Plasmin", J. Med. Chem., 43:3093-3102 (2000).	

EXAMINER /Nizal Chandrakumar/

DATE CONSIDERED

09/06/2009



**INFORMATION DISCLOSURE  
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Form PTO-1449 (Modified)  
(Use several sheets if necessary)

Sheet 2 of 2

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17. /N.C./	Dittert, L.W. et al., "Acetaminophen Prodrugs I Synthesis, Physicochemical Properties, and Analgesic Activity", <i>Journal of Pharmaceutical Sciences</i> , 57(5):774-780 (1968).
18.	Dittert, L.W. et al., "Acetaminophen Prodrugs II Effect of Structure and Enzyme Source on Enzymatic and Nonenzymatic Hydrolysis of Carbonate Esters," <i>J. of Pharm. Sciences</i> , 57(5):780-783 (1968).
19.	Hansen et al., "Carbamate Ester Prodrugs of Dopaminergic Compounds: Synthesis, Stability, and Bioconversion", <i>Journal of Pharmaceutical Sciences</i> , 80(8):793-798 (1991).
20.	Hansen et al., "Ketobemidone prodrugs for buccal delivery", <i>Acta Pharm. Nord.</i> , 3(2):77-82 (1991).
21.	Huang et al., "Hydrolysis of Carbonates, Thiocarbonates, Carbamates, and Carboxylic Esters of α-Naphthol, β-Naphthol, and p-Nitrophenol by Human, Rat, and Mouse Liver Carboxylesterases", <i>Pharmaceutical Research</i> , 10(5):639-648 (1993).
22.	Kahns et al., "Prodrugs of Peptides. 18. Synthesis and Evaluation of Various Esters of Desmopressin (dDAVP)", <i>Pharmaceutical Research</i> , 10(1):68-74 (1993).
23.	Nassar et al., "Effects of Structural Variations on the Rates of Enzymatic and Nonenzymatic Hydrolysis of Carbonate and Carbamate Esters", <i>Journal of Pharmaceutical Sciences</i> , 81(3):295-298 (1992).
24.	Savolainen et al., "Synthesis and <i>in vitro</i> <i>in vivo</i> evaluation of novel oral N-alkyl- and <i>N,N</i> -dialkyl-carbamate esters of entacapone," <i>Life Sciences</i> , 67:205-216 (2000).
25.	Tunek et al., "Hydrolysis of <sup>3</sup> H-Bambuterol, A Carbamate Prodrug of Terbutaline, in Blood from Humans and Laboratory Animals <i>In Vitro</i> ", <i>Biochemical Pharmacology</i> , 37(20):3867-3876 (1988).
26.	Weibel et al., "Macromolecular prodrugs IX. Kinetics of hydrolysis of benzyl dextran carbonate ester conjugates in aqueous buffer solutions and human plasma", <i>Acta Pharm. Nord.</i> , 3(3):159-162 (1991).
27.	Yang et al., <i>Tetrahedron Letters</i> , 38(39):6865-6868 (1997).
28. /N.C./	Yu et al., <i>Chinese Chemical Letters</i> , 2(12):937-940 (1991).

EXAMINER	DATE CONSIDERED
/Nizal Chandrakumar/	09/06/2009